EAST Search History

Ref #	Hits	Search Query	DBs	Default Operator	Plurals	Time Stamp
L1	3	"7265123"	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/11/27 09:49
L2	0	us2006216288	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/11/27 09:49
L3	2	"2006216288"	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/11/27 09:50
L4	2	"20060216288"	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/11/27 09:50
L6	8608	"Spector".in. or "Xia".in.	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/11/27 09:52
L7	2	l6 and (craf-1 or craf1)	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/11/27 09:53
L8	0	(craf-1 or craf1) same combin?	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/11/27 09:53
L9	119	(craf-1 or craf1) same erbb2	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/11/27 09:53

EAST Search History

S1	16	"6268391"	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/11/27 09:48
S2	3	"7084147"	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/04/16 10:27
S3	5	"6719339"	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/04/16 10:27
S4	3	"7109333"	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/04/16 10:50
S5	9	"6727256"	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/04/16 14:04
S6	2	"7189734"	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/11/27 09:51
S7	2	"7141576"	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/04/16 11:11
S8	12	"6713485"	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/04/16 11:11

EAST Search History

S9	7	"bRaf inhibitor"	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/04/16 14:38
S10	64	"Raf inhibitor"	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/04/16 14:38
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S12	10	"b-Raf inhibitor" and "cancer"	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/04/16 15:33
S13	0	514/264.110	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/04/16 15:34
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S16	166	514/264.11.ccls.	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/04/16 15:34

EAST Search History

S17	22	514/264.11.ccls. and ("erbb2" or "raf")	US-PGPUB; USPAT; USOCR; FPRS; EPO; JPO; DERWENT	OR	OFF	2007/04/16 15:35

10/510,542
11/27/07

Connecting via Winsock to STN

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NEWS	2	JUL 02	LMEDLINE coverage updated
NEWS	3	JUL 02	SCISEARCH enhanced with complete author names
NEWS	4	JUL 02	CHEMCATS accession numbers revised
NEWS	5	JUL 02	CA/CAplus enhanced with utility model patents from China
NEWS	6	JUL 16	CAplus enhanced with French and German abstracts
NEWS	7	JUL 18	CA/CAplus patent coverage enhanced
NEWS	8	JUL 26	USPATFULL/USPAT2 enhanced with IPC reclassification
NEWS	9	JUL 30	USGENE now available on STN
NEWS	10	AUG 06	CAS REGISTRY enhanced with new experimental property tags
NEWS	11	AUG 06	FSTA enhanced with new thesaurus edition
NEWS	12	AUG 13	CA/CAplus enhanced with additional kind codes for granted patents
NEWS	13	AUG 20	CA/CAplus enhanced with CAS indexing in pre-1907 records
NEWS	14	AUG 27	Full-text patent databases enhanced with predefined patent family display formats from INPADOCDB
NEWS	15	AUG 27	USPATOLD now available on STN
NEWS	16	AUG 28	CAS REGISTRY enhanced with additional experimental spectral property data
NEWS	17	SEP 07	STN AnaVist, Version 2.0, now available with Derwent World Patents Index
NEWS	18	SEP 13	FORIS renamed to SOFIS
NEWS	19	SEP 13	INPADOCDB enhanced with monthly SDI frequency
NEWS	20	SEP 17	CA/CAplus enhanced with printed CA page images from 1967-1998
NEWS	21	SEP 17	CAplus coverage extended to include traditional medicine patents
NEWS	22	SEP 24	EMBASE, EMBAL, and LEMBASE reloaded with enhancements
NEWS	23	OCT 02	CA/CAplus enhanced with pre-1907 records from Chemisches Zentralblatt
NEWS	24	OCT 19	BEILSTEIN updated with new compounds
NEWS	25	NOV 15	Derwent Indian patent publication number format enhanced
NEWS	26	NOV 19	WPIX enhanced with XML display format

NEWS EXPRESS 19 SEPTEMBER 2007: CURRENT WINDOWS VERSION IS V8.2, CURRENT MACINTOSH VERSION IS V6.0c(ENG) AND V6.0Jc(JP), AND CURRENT DISCOVER FILE IS DATED 19 SEPTEMBER 2007.

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FILE 'HOME' ENTERED AT 09:44:02 ON 27 NOV 2007

=> file registry

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	0.21	0.21

FILE 'REGISTRY' ENTERED AT 09:44:14 ON 27 NOV 2007
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STRUCTURE FILE UPDATES: 26 NOV 2007 HIGHEST RN 955995-34-3
DICTIONARY FILE UPDATES: 26 NOV 2007 HIGHEST RN 955995-34-3

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on property searching in REGISTRY, refer to:

http://www.cas.org/support/stngen/stndoc/properties.html

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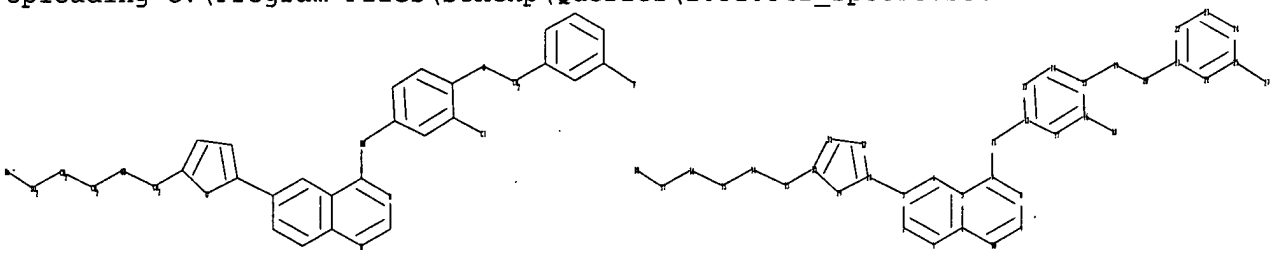

chain nodes :
11 18 19 20 27 33 34 35 36 37 38
ring nodes :
1 2 3 4 5 6 7 8 9 10 12 13 14 15 16 17 21 22 23 24 25 26 28
29 30 31 32
chain bonds :
3-28 7-11 11-12 15-19 16-18 19-20 20-21 25-27 30-33 33-34 34-35 35-36
36-37 37-38

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ring bonds :
1-2  1-6  2-3  3-4  4-5  5-6  5-7  6-10  7-8  8-9  9-10  12-13  12-17  13-14  14-15
 15-16  16-17  21-22  21-26  22-23  23-24  24-25  25-26  28-29  28-32  29-30  30-31
 31-32
exact/norm bonds :
7-11  11-12  15-19  28-29  28-32  29-30  30-31  31-32
exact bonds :
3-28  16-18  19-20  20-21  25-27  30-33  33-34  34-35  35-36  36-37  37-38
normalized bonds :
1-2  1-6  2-3  3-4  4-5  5-6  5-7  6-10  7-8  8-9  9-10  12-13  12-17  13-14  14-15
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Match level :
1:Atom  2:Atom  3:Atom  4:Atom  5:Atom  6:Atom  7:Atom  8:Atom  9:Atom  10:Atom
11:CLASS  12:Atom  13:Atom  14:Atom  15:Atom  16:Atom  17:Atom  18:CLASS  19:CLASS
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37:CLASS  38:CLASS



L1      STRUCTURE UPLOADED

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L1 HAS NO ANSWERS
L1              STR
*** STRUCTURE DIAGRAM IS NOT AVAILABLE ***

Structure attributes must be viewed using STN Express query preparation.


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FULL SEARCH INITIATED 09:44:41 FILE 'REGISTRY'
FULL SCREEN SEARCH COMPLETED -       26 TO ITERATE

100.0% PROCESSED       26 ITERATIONS                                  1 ANSWERS
SEARCH TIME: 00.00.01

L2            1 SEA EXA FUL L1

=> d l2

L2   ANSWER 1 OF 1  REGISTRY  COPYRIGHT 2007 ACS on STN
RN   231277-92-2  REGISTRY
ED   Entered STN:  07 Aug 1999
CN   4-Quinazolinamine, N-[3-chloro-4-[(3-fluorophenyl)methoxy]phenyl]-6-[5-
     [[[2-(methylsulfonyl)ethyl]amino]methyl]-2-furanyl]-  (CA INDEX NAME)
OTHER NAMES:
CN   4-[[3-Chloro-4-(3-fluorobenzyloxy)phenyl]amino]-6-[5-[[(2-
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CN   GSK 572016
CN   GW 572016
CN   Lapatinib
MF   C29 H26 Cl F N4 O4 S
CI   COM
SR   CA
LC   STN Files:   ADISINSIGHT, ANABSTR, BIOSIS, CA, CAPLUS, CASREACT, CBNB,
       CHEMCATS, EMBASE, IMSDRUGNEWS, IMSPATENTS, IMSRESEARCH, IPA, PROUSDDR,
       RTECS*, SYNTHLINE, TOXCENTER, USAN, USPAT2, USPATFULL
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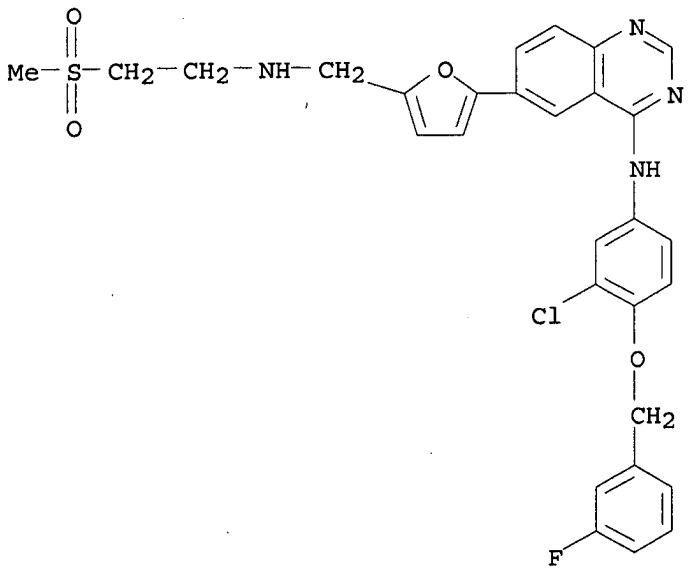

PROPERTY DATA AVAILABLE IN THE 'PROP' FORMAT

216 REFERENCES IN FILE CA (1907 TO DATE)
4 REFERENCES TO NON-SPECIFIC DERIVATIVES IN FILE CA
218 REFERENCES IN FILE CAPLUS (1907 TO DATE)

=> file medline caplus wpids uspatfull

COST IN U.S. DOLLARS	SINCE FILE ENTRY	TOTAL SESSION
FULL ESTIMATED COST	60.20	60.41

FILE 'MEDLINE' ENTERED AT 09:45:00 ON 27 NOV 2007

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FILE 'USPATFULL' ENTERED AT 09:45:00 ON 27 NOV 2007
CA INDEXING COPYRIGHT (C) 2007 AMERICAN CHEMICAL SOCIETY (ACS)

=> s l2
SAMPLE SEARCH INITIATED 09:45:04 FILE 'WPIDS'
SAMPLE SCREEN SEARCH COMPLETED - 3 TO ITERATE

100.0% PROCESSED 3 ITERATIONS 1 ANSWERS
SEARCH TIME: 00.00.01

FULL FILE PROJECTIONS:	ONLINE	**COMPLETE**
	BATCH	**COMPLETE**
PROJECTED ITERATIONS:	3 TO	81
PROJECTED ANSWERS:	1 TO	40

L3 272 L2

=> s l3 and (craf-1 or craf1)
L4 6 L3 AND (CRAF-1 OR CRAF1)

=> d l4 1-6 ibib, abs, hitstr

L4 ANSWER 1 OF 6 CAPLUS COPYRIGHT 2007 ACS on STN
ACCESSION NUMBER: 2003:836903 CAPLUS
DOCUMENT NUMBER: 139:317433
TITLE: Cancer treatment method comprising administering an erb-family inhibitor and a raf and/or ras inhibitor
INVENTOR(S): Spector, Neil Lee; Xia, Wenle
PATENT ASSIGNEE(S): Smithkline Beecham Corporation, USA
SOURCE: PCT Int. Appl., 173 pp.
CODEN: PIXXD2
DOCUMENT TYPE: Patent
LANGUAGE: English
FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2003086467	A1	20031023	WO 2003-US10747	20030408
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NI, NO, NZ, OM, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW				
RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
AU 2003221684	A1	20031027	AU 2003-221684	20030408
EP 1492568	A1	20050105	EP 2003-718262	20030408
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, LT, LV, FI, RO, MK, CY, AL, TR, BG, CZ, EE, HU, SK				
JP 2005534623	T	20051117	JP 2003-583483	20030408
US 2005176740	A1	20050811	US 2004-510542	20041007
PRIORITY APPLN. INFO.:			US 2002-370807P	P 20020408
			WO 2003-US10747	W 20030408

OTHER SOURCE(S): MARPAT 139:317433
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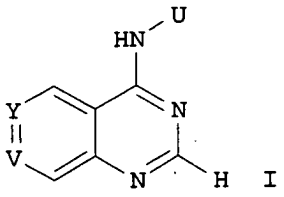


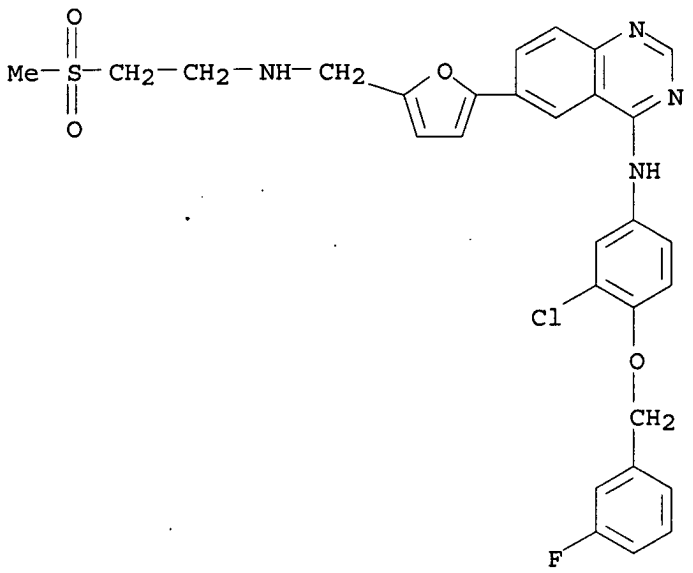
I

AB The invention provides a method for treating cancer in a mammal, as well as pharmaceutical combinations useful in such treatment. In particular, the method relates to a cancer treatment method that includes administering an erb family inhibitor and a Raf and/or ras inhibitor to a mammal suffering from a cancer. Preparation of compds., e.g. erbB-2/EGFR inhibitor I, is described.

IT 231277-92-2P
RL: PAC (Pharmacological activity); SPN (Synthetic preparation); THU (Therapeutic use); BIOL (Biological study); PREP (Preparation); USES (Uses)
(erb-family inhibitor and raf and/or ras inhibitor combination for cancer treatment)

RN 231277-92-2 CAPLUS

CN 4-Quinazolinamine, N-[3-chloro-4-[(3-fluorophenyl)methoxy]phenyl]-6-[5-[[[2-(methylsulfonyl)ethyl]amino]methyl]-2-furanyl]- (CA INDEX NAME)



REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

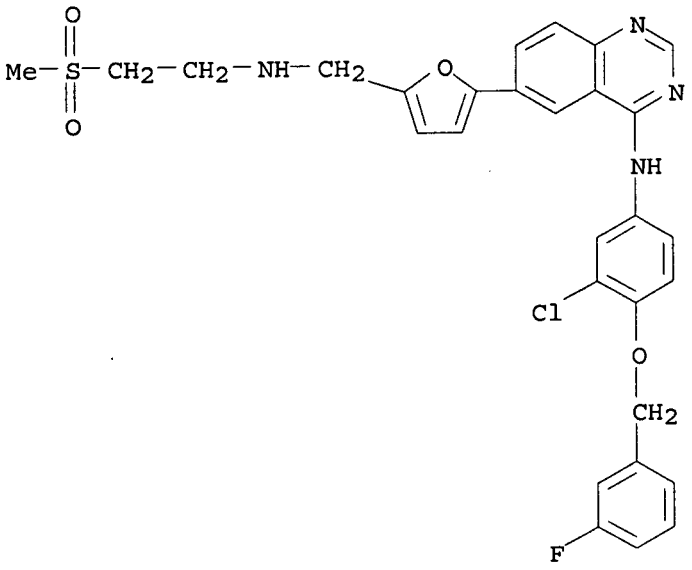
L4 ANSWER 2 OF 6 USPATFULL on STN

ACCESSION NUMBER:	2007:107539 USPATFULL		
TITLE:	HETEROCYCLIC COMPOUNDS		
INVENTOR(S):	COCKERILL, George Stuart, Maulden, UNITED KINGDOM Lackey, Karen Elizabeth, Durham, NC, UNITED STATES		
	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2007093512	A1	20070426
	US 7265123	B2	20070904
APPLICATION INFO.:	US 2006-562047	A1	20061121 (11)
RELATED APPLN. INFO.:	Division of Ser. No. US 2006-400284, filed on 7 Apr 2006, GRANTED, Pat. No. US 7189734 Division of Ser. No. US 2005-61578, filed on 18 Feb 2005, GRANTED, Pat. No. US 7084147 Division of Ser. No. US 2002-30527, filed on 9 Jan 2002, GRANTED, Pat. No. US 6933299		
	NUMBER	DATE	
PRIORITY INFORMATION:	GB 1999-16213	19990709	
	GB 1999-16218	19990709	
DOCUMENT TYPE:	Utility		
FILE SEGMENT:	APPLICATION		
LEGAL REPRESENTATIVE:	GLAXOSMITHKLINE, CORPORATE INTELLECTUAL PROPERTY, MAI B475, FIVE MOORE DR., PO BOX 13398, RESEARCH TRIANGLE PARK, NC, 27709-3398, US		
NUMBER OF CLAIMS:	29		
EXEMPLARY CLAIM:	1		
LINE COUNT:	4376		

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Heteroaromatic compounds are described, methods for their preparation, pharmaceutical compositions containing them, methods of use, and their use in medicines. In particular, the invention relates to quinazoline and pyridopyrimidine derivatives which exhibit protein tyrosine kinase inhibition.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.
IT 231277-92-2
(preparation of anilinoquinazolines as protein tyrosine kinase inhibitors)
RN 231277-92-2 USPATFULL
CN 4-Quinazolinamine, N-[3-chloro-4-[(3-fluorophenyl)methoxy]phenyl]-6-[5-[[[2-(methylsulfonyl)ethyl]amino]methyl]-2-furanyl]- (CA INDEX NAME)



L4 ANSWER 3 OF 6 USPATFULL on STN

ACCESSION NUMBER:	2006:253838 USPATFULL
TITLE:	Combinations for the treatment of cancer
INVENTOR(S):	Chang, David, Calabasas, CA, UNITED STATES
PATENT ASSIGNEE(S):	Amgen Inc, Thousand Oaks, CA, UNITED STATES (U.S. corporation)

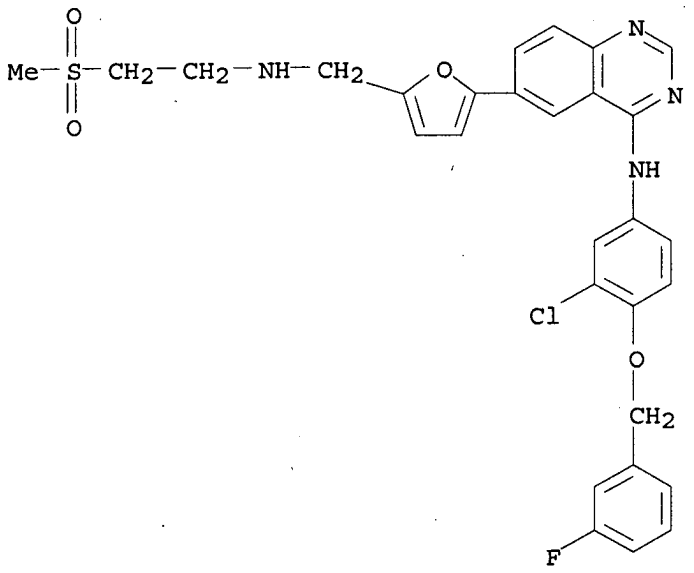
	NUMBER	KIND	DATE	
PATENT INFORMATION:	US 2006216288	A1	20060928	
APPLICATION INFO.:	US 2006-386271	A1	20060321	(11)

	NUMBER	DATE	
PRIORITY INFORMATION:	US 2005-664381P	20050322	(60)

DOCUMENT TYPE:	Utility
FILE SEGMENT:	APPLICATION
LEGAL REPRESENTATIVE:	AMGEN INC., MAIL STOP 28-2-C, ONE AMGEN CENTER DRIVE, THOUSAND OAKS, CA, 91320-1799, US
NUMBER OF CLAIMS:	15
EXEMPLARY CLAIM:	1
NUMBER OF DRAWINGS:	5 Drawing Page(s)
LINE COUNT:	1584

CAS INDEXING IS AVAILABLE FOR THIS PATENT.
AB This invention is in the field of pharmaceutical agents and specifically relates to compounds, compositions, uses and methods for treating cancer.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.
IT 231277-92-2
(combinations for the treatment of cancer)
RN 231277-92-2 USPATFULL
CN 4-Quinazolinamine, N-[3-chloro-4-[(3-fluorophenyl)methoxy]phenyl]-6-[5-[[[2-(methylsulfonyl)ethyl]amino]methyl]-2-furanyl]- (CA INDEX NAME)



L4 ANSWER 4 OF 6 USPATFULL on STN

ACCESSION NUMBER:	2006:222351 USPATFULL
TITLE:	Anilinoquinazaolines as protein tyrosine kianse inhibitors
INVENTOR(S):	Cockerill, George Stuart, Maulden, UNITED KINGDOM Lackey, Karen Elizabeth, Durham, NC, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2006189637	A1	20060824
	US 7189734	B2	20070313
APPLICATION INFO.:	US 2006-400284	A1	20060407 (11)

RELATED APPLN. INFO.: Division of Ser. No. US 2005-61578, filed on 18 Feb 2005, PENDING Division of Ser. No. US 2002-30527, filed on 9 Jan 2002, GRANTED, Pat. No. US 6933299

	NUMBER	DATE
PRIORITY INFORMATION:	GB 1999-16213	19990709
	GB 1999-16218	19990709

DOCUMENT TYPE:	Utility
FILE SEGMENT:	APPLICATION
LEGAL REPRESENTATIVE:	GLAXOSMITHKLINE, CORPORATE INTELLECTUAL PROPERTY, MAI B475, FIVE MOORE DR., PO BOX 13398, RESEARCH TRIANGLE PARK, NC, 27709-3398, US
NUMBER OF CLAIMS:	25
EXEMPLARY CLAIM:	1
LINE COUNT:	4471

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

AB Heteroaromatic compounds are described, methods for their preparation, pharmaceutical compositions containing them, methods of use, and their use in medicines. In particular, the invention relates to quinazoline and pyridopyrimidine derivatives which exhibit protein tyrosine kinase inhibition.

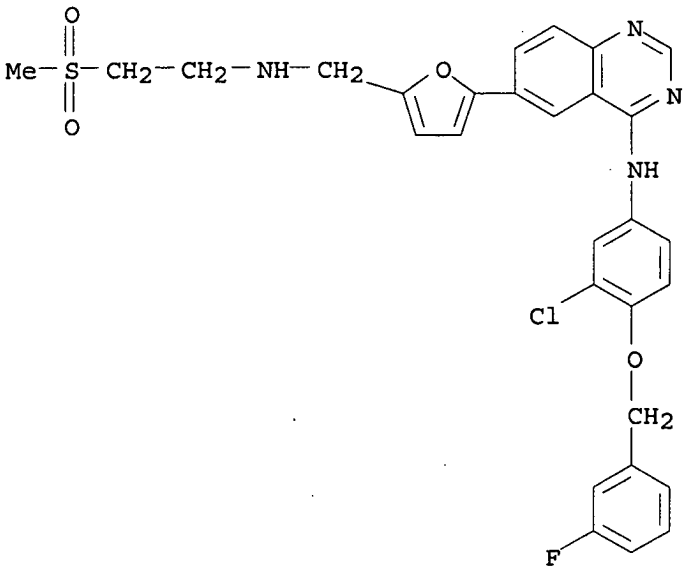
CAS INDEXING IS AVAILABLE FOR THIS PATENT.

IT 231277-92-2
(preparation of anilinoquinazolines as protein tyrosine kinase inhibitors)

RN 231277-92-2 USPATFULL

CN 4-Quinazolinamine, N-[3-chloro-4-[(3-fluorophenyl)methoxy]phenyl]-6-[5-

[[[2-(methylsulfonyl)ethyl]amino]methyl]-2-furanyl]- (CA INDEX NAME)



L4 ANSWER 5 OF 6 USPATFULL on STN

ACCESSION NUMBER:	2005:203315 USPATFULL
TITLE:	Cancer treatment method comprising administering an erb-family inhibitor and a raf and/or ras inhibitor
INVENTOR(S):	Spector, Neil Lee, Durham, NC, UNITED STATES Xia, Wenle, Durham, NC, UNITED STATES

	NUMBER	KIND	DATE	
PATENT INFORMATION:	US 2005176740	A1	20050811	
APPLICATION INFO.:	US 2003-510542	A1	20030408	(10)
	WO 2003-US10747		20030408	

	NUMBER	DATE
PRIORITY INFORMATION:	US 2002-370807P	20020408 (60)

DOCUMENT TYPE:	Utility
FILE SEGMENT:	APPLICATION
LEGAL REPRESENTATIVE:	DAVID J LEVY, CORPORATE INTELLECTUAL PROPERTY, GLAXOSMITHKLINE, FIVE MOORE DR., PO BOX 13398, RESEARCH TRIANGLE PARK, NC, 27709-3398, US
NUMBER OF CLAIMS:	25
EXEMPLARY CLAIM:	1
NUMBER OF DRAWINGS:	10 Drawing Page(s)
LINE COUNT:	3918

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

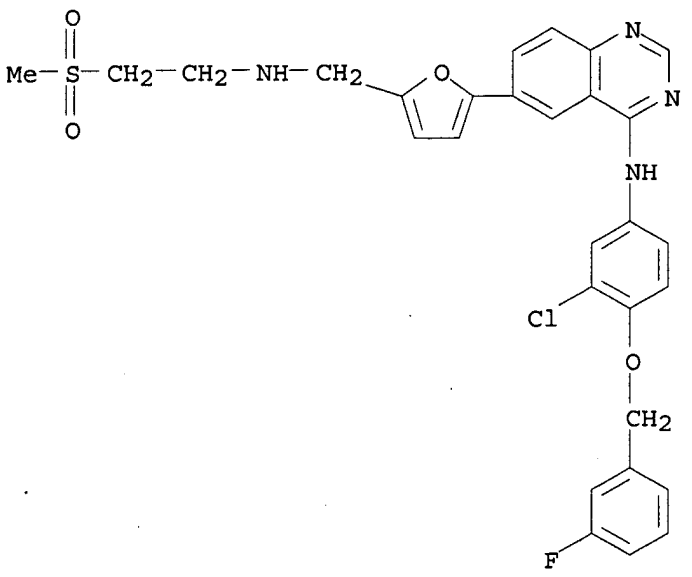
AB The present invention relates to a method of treating cancer in a mammal and to pharmaceutical combinations useful in such treatment. In particular, the method relates to a cancer treatment method that includes administering an erb family inhibitor and a Raf and/or ras inhibitor to a mammal suffering from a cancer.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

IT 231277-92-2P
(erb-family inhibitor and raf and/or ras inhibitor combination for cancer treatment)

RN 231277-92-2 USPATFULL

CN 4-Quinazolinamine, N-[3-chloro-4-[(3-fluorophenyl)methoxy]phenyl]-6-[5-[[[2-(methylsulfonyl)ethyl]amino]methyl]-2-furanyl]- (CA INDEX NAME)



L4 ANSWER 6 OF 6 USPATFULL on STN

ACCESSION NUMBER: 2005:165976 USPATFULL

TITLE: Anilinoquinazaolines as protein tyrosine kianse inhibitors

INVENTOR(S): Cockerill, George Stuart, Maulden, UNITED KINGDOM
Lackey, Karen Elizabeth, Durham, NC, UNITED STATES

	NUMBER	KIND	DATE
PATENT INFORMATION:	US 2005143401	A1	20050630
	US 7084147	B2	20060801
APPLICATION INFO.:	US 2005-61578	A1	20050218 (11)

RELATED APPLN. INFO.: Division of Ser. No. US 2002-303527, filed on 25 Nov 2002, GRANTED, Pat. No. US 6719339

	NUMBER	DATE
PRIORITY INFORMATION:	GB 1999-16213	19990709
	GB 1999-16218	19990709

DOCUMENT TYPE: Utility

FILE SEGMENT: APPLICATION

LEGAL REPRESENTATIVE: DAVID J LEVY, CORPORATE INTELLECTUAL PROPERTY, GLAXOSMITHKLINE, FIVE MOORE DR., PO BOX 13398, RESEARCH TRIANGLE PARK, NC, 27709-3398, US

NUMBER OF CLAIMS: 32

EXEMPLARY CLAIM: 1

LINE COUNT: 4418

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

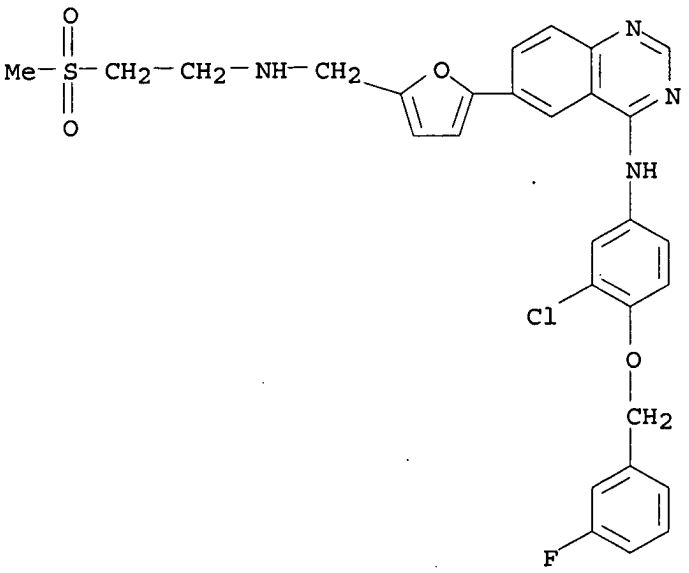
AB Heteroaromatic compounds are described, methods for their preparation, pharmaceutical compositions containing them, methods of use, and their use in medicines. In particular, the invention relates to quinazoline and pyridopyrimidine derivatives which exhibit protein tyrosine kinase inhibition.

CAS INDEXING IS AVAILABLE FOR THIS PATENT.

IT 231277-92-2
(preparation of anilinoquinazolines as protein tyrosine kinase inhibitors)

RN 231277-92-2 USPATFULL

CN 4-Quinazolinamine, N-[3-chloro-4-[(3-fluorophenyl)methoxy]phenyl]-6-[5-[[[2-(methylsulfonyl)ethyl]amino]methyl]-2-furanyl]- (CA INDEX NAME)



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     FILE 'REGISTRY' ENTERED AT 09:44:14 ON 27 NOV 2007
L1              STRUCTURE UPLOADED
L2            1 S L1 EXA FULL

     FILE 'MEDLINE, CAPLUS, WPIDS, USPATFULL' ENTERED AT 09:45:00 ON 27 NOV
     2007
L3          272 S L2
L4            6 S L3 AND (CRAF-1 OR CRAF1)

=>

---Logging off of STN---


=>
Executing the logoff script...


=> LOG Y

COST IN U.S. DOLLARS                                SINCE FILE      TOTAL
                                                       ENTRY    SESSION
FULL ESTIMATED COST                                    47.63     108.04

DISCOUNT AMOUNTS (FOR QUALIFYING ACCOUNTS)          SINCE FILE      TOTAL
                                                       ENTRY    SESSION
CA SUBSCRIBER PRICE                                    -0.78      -0.78

STN INTERNATIONAL LOGOFF AT 09:46:27 ON 27 NOV 2007
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